

Maryland Lyme Disease (LD) Case Report Form

PATIENT INFORMATION																																																															
NAME OF PATIENT – LAST FIRST M				TELEPHONE NUMBERS Home: Workplace:		DATE reported to HD: NEDSS: CAS																																																									
Address:				Zip Code:		County of Residence:																																																									
Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Unknown		Date of birth: (mmddyyyy)		Hispanic Ethnicity: <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown		Race: <input type="checkbox"/> American Indian or Alaskan Native <input type="checkbox"/> Asian <input type="checkbox"/> Black or African American <input type="checkbox"/> White <input type="checkbox"/> Native Hawaiian or Pacific Islander <input type="checkbox"/> Unknown																																																									
PHYSICIAN / PROVIDER INFORMATION																																																															
Physician:				Phone:		FAX:																																																									
LABORATORY FINDINGS																																																															
EIA/IFA (IgM and/or IgG) <input type="checkbox"/> positive <input type="checkbox"/> equivocal <input type="checkbox"/> negative <input type="checkbox"/> not done <input type="checkbox"/> check if assay uses C6 peptide Specimen collection date: (if not serum, specify):																																																															
Western Blot (WB) IgM: <input type="checkbox"/> positive <input type="checkbox"/> negative <input type="checkbox"/> not done Specimen collection date: <input type="checkbox"/> 41kDa (FlaB) <input type="checkbox"/> 39 kDa (BmpA) <input type="checkbox"/> 21-25 kDa (OspC) (if not serum, specify): IgG: <input type="checkbox"/> positive <input type="checkbox"/> negative <input type="checkbox"/> not done Please indicate <u>positive WB bands</u> , if known. For IgM, 2 of 3 bands must be positive <input type="checkbox"/> 93 kDa <input type="checkbox"/> 66 kDa <input type="checkbox"/> 58 kDa <input type="checkbox"/> 45 kDa <input type="checkbox"/> 41 kDa For IgG, 5 of 10 bands must be positive <input type="checkbox"/> 39 kDa <input type="checkbox"/> 30 kDa <input type="checkbox"/> 28 kDa <input type="checkbox"/> 21 kDa <input type="checkbox"/> 18 kDa																																																															
Other tests (check what applies): <input type="checkbox"/> <i>B. burgdorferi</i> cultured <input type="checkbox"/> CSF titer higher than serum titer*? <input type="checkbox"/> Other (please specify): Specimen collection date:																																																															
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LYME DISEASE (LD) SURVEILLANCE CASE DEFINITION (07-ID-11)

Clinical description:

A systemic, tickborne disease with protean manifestations, including dermatologic, rheumatologic, neurologic, and cardiac abnormalities. The best clinical marker for the disease is the initial skin lesion (i.e., erythema migrans {EM}) that occurs in 60%-80% of patients.

Surveillance case definition:

This surveillance case definition was developed for national reporting of Lyme disease; it is not intended to be used in clinical diagnosis.

Case classifications:

Confirmed case:

- EM with a known exposure (as defined below), *or*
- EM with laboratory evidence (as defined below) of infection and without a known exposure, *or*
- At least one late manifestation that has laboratory evidence of infection

Probable:

- Physician-diagnosed LD that has laboratory evidence of infection with non-confirmatory* signs and symptoms

Suspect:

- A case of EM where there is no known exposure and no evidence of infection, *or*
- A case with laboratory evidence of infection but no clinical information available (e.g. a laboratory report)

Definitions and Clarifications:

Erythema migrans (EM). For purposes of surveillance, EM is defined as a skin lesion that typically begins as a red macule or papule and expands over a period of days to weeks to form a large round lesion, often with partial central clearing. A single primary lesion must reach greater than or equal to 5 cm in size. Secondary lesions also may occur. Annular erythematous lesions occurring within several hours of a tick bite represent hypersensitivity reactions and do not qualify as EM. For most patients, the expanding EM lesion is accompanied by other acute symptoms, particularly fatigue, fever, headache, mildly stiff neck, arthralgia, or myalgia. These symptoms are typically intermittent.

The diagnosis of EM must be made by a physician. Laboratory confirmation is recommended for persons with no known exposure.

Confirmatory late manifestations include any of the following when an alternate explanation is not found:

1. Musculoskeletal system. Recurrent, brief attacks (weeks or months) of objective joint swelling in one or a few joints, sometimes followed by chronic arthritis in one or a few joints. Manifestations not considered as criteria for diagnosis include chronic progressive arthritis not preceded by brief attacks and chronic symmetrical polyarthritis.
2. Nervous system. Any of the following, alone or in combination: lymphocytic meningitis; cranial neuritis, particularly facial palsy (may be bilateral); radiculoneuropathy; or, rarely, encephalomyelitis. Encephalomyelitis must be confirmed by demonstration of antibody production against *B. burgdorferi* in the CSF, evidenced by a higher titer of antibody in CSF than in serum.
3. Cardiovascular system. Acute onset of high-grade (2nd-degree or 3rd-degree) atrioventricular conduction defects that resolve in days to weeks and are sometimes associated with myocarditis.

***Non-confirmatory.** Non-confirmatory signs and symptoms include:

Fever, sweats, chills, fatigue, neck pain, arthralgias, myalgias, fibromyalgia syndromes, cognitive impairment, headache, paresthesias, visual/auditory impairment, peripheral neuropathy, encephalopathy, palpitations, bradycardia, bundle branch block, myocarditis, or other rash.

Exposure. Exposure is defined as having been (≤ 30 days before onset of EM) in wooded, brushy, or grassy areas (i.e., potential tick habitats) in a county in which Lyme disease is endemic. A history of tick bite is not required.

Disease endemic to county. A county in which Lyme disease is endemic in which at least two confirmed cases have been acquired in the county or in which established populations of a known tick vector are infected with *B. burgdorferi*.

Laboratory evidence. For the purpose of surveillance, the definition of a qualified laboratory assay is

1. A positive culture for *B. burgdorferi*,
2. Two-tier testing with IgM or IgG immunoblot seropositive interpreted using established criteria.
Note: A positive IgM test result alone is not recommended for use in determining active disease in persons with illness greater than 1 month's duration because the likelihood of a false-positive test result for a current infection is high in these persons.
3. Single-tier IgG immunoblot seropositive interpreted using established criteria. Additional assays may be added based on periodic review of the scientific literature and strong evidence of comparable or better performance than qualifying assays.